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WE, Shanta Modak, Trupti A. Gaonkar, and Lester Sampath, citizens of the United States, India, and the United States, respectively, having post office addresses at 184 Howland Avenue, Riveredge, New Jersey 07661; 712 W. 176th Street, Apt. #5F, New York, NY 10033; and 7 Lawrence Street, Nyack, New York 10960, respectively, have made an invention in

GENTLE-ACTING SKIN-DISINFECTANTS

of which the following is a

SPECIFICATION

1. INTRODUCTION

[0001] The present invention provides for skin-friendly antimicrobial compositions comprising synergistic combinations of octoxyglycerin and a low concentration of an antibiotic, particularly chlorhexidine. In particular embodiments, the compositions further comprise a quaternary ammonium compound that enhances killing of microbes.

2. BACKGROUND OF THE INVENTION

[0002] "Skin disinfectants" are routinely used in professional and non-professional contexts to rapidly kill microbes. A physician has a need to disinfect his or her skin both before and after examining a patient. Prior to the performance of an invasive medical procedure, the skin of the subject must be properly cleaned to avoid post-procedure infections. In non-professional contexts, a commuter, riding public transportation, may wish to disinfect her hands before handling food; a child, playing in a park, may need to clean his hands but not have the convenience of soap and water nearby. Each of these situations require, optimally, a skin disinfectant that is effective, easy to use, and non-irritating so as to permit repeated use.

[0003] A number of skin disinfectants have been developed that use alcohol as the primary antimicrobial agent. There are two general problems associated with alcohol-based disinfectants. First, the effective concentration of alcohol, generally regarded to be greater than about 60 percent weight (hereafter, all percentages should be considered weight/volume percentages, unless specified otherwise) of ethanol, or its equivalent, is irritating to the skin, causing dryness and consequent peeling and cracking. Because chapped skin tends to be more susceptible to microbial contamination, repeated use of alcohol disinfectants can exacerbate the very problem they are intended to solve. Second, whereas alcohol can be an effective disinfectant, once it evaporates its antimicrobial activity is lost.

[0004] Alcohol-based skin disinfectants which are known in the art, some of which address the two problems mentioned above, include the following.

[0005] United States Patent No. 6,107, 261 by Taylor et al., issued August 22, 2000, and its continuations-in-part, United States Patent No. 6,204,230 by Taylor et al., issued March 20, 2001 and United States Patent No. 6,136,771 by Taylor et al., issued October 24, 2000, disclose antibacterial compositions which contain an antibacterial agent at a percent saturation of at least 50 percent. The compositions further comprise, as solubility promoters, a surfactant and a hydric solvent, which may be an alcohol.

[0006] United States Patent No. 5,776,430 by Osborne et al., issued July 7, 1998, discloses a topical antimicrobial cleaner containing about 0.65 -0.85 percent chlorhexidine and about 50-60 percent denatured alcohol, which is scrubbed onto and then rinsed off the skin.

[0007] European Patent Application 0604 848 discloses a gel comprising an antimicrobial agent, 40-90 percent by weight of an alcohol, and a polymer and thickening agent.

[0008] United States Patent No. 4,956,170 by Lee, issued September 11, 1990 relates to a high alcohol content antimicrobial gel composition which comprises various emollients and a humectant to protect the skin from the drying effects of the alcohol. In alcohol formulations, higher levels of alcohol are needed to provide instant kill against sensitive as well as resistant strains of bacteria.

[0009] Certain formulations virtually omit alcohol as a primary antimicrobial agent, such as, for example, the skin sanitizing compositions disclosed in United States Patent No. 6,187,327 by Stack, issued February 13, 2001, which comprises triclosan (2,4,4'-trichloro-2'-hydroxydiphenyl ether; concentration 0.1-0.35 weight percent) in a topical lotion comprised of a surfactant phase and a wax phase, which purportedly provides antimicrobial protection for 3-4 hours after application. The composition prepared according to the claims of United States Patent No. 6,187,327 further comprises chlorhexidine digluconate.

[0010] United States Patent No. 5,965,610 by Modak et al., issued October 12, 1999, teaches skin cleaning compositions comprising antimicrobial agents and zinc salts, where zinc salts have a soothing effect on the skin. The claimed subject matter includes formulations comprising a gel formed between zinc gluconate, chlorhexidine gluconate and a solvent, to which various thickening agents, emulsifying agents and/or emollients may be added.

[0011] United States Patent No. 5,985,918 by Modak et al., issued November 16, 1999, relates to "Zinc-Based Anti-Irritant Creams".

[0012] United States Patent No. 5,705,532 by Modak et al., issued January 6, 1998, relates to "Triple Antimicrobial Compositions" comprising less than or equal to two percent of a chlorhexidine compound, less than or equal to 0.1 percent of a quaternary ammonium compound, and less than or equal to two percent parachlorometaxlenol.

[0013] Octoxyglycerin, sold under the trade name Sensiva® SC50 (Schulke & Mayr), is a glycerol alkyl ether known to be gentle to the skin. Octoxyglycerine exhibits antimicrobial activity against a variety of Gram-positive bacteria associated with perspiration odor, such as *Micrococcus luteus*, *Corynebacterium aquaticum*, *Corynebacterium flavescent*, *Corynebacterium callunae*, and *Corynebacterium nephredi*, and is used in various skin deodorant preparations at concentrations between about 0.2 and 3 percent (Sensiva® product literature, Schulke & Mayr).

[0014] For example, United States Patent No. 5,885,562 by Lowry et al., issued March 23, 1999, relates to deodorant compositions comprising an antimicrobial agent, namely polyhexamethylene biguanide (at a concentration of between 0.01 and 0.5 percent), together with a polarity modifier such as Sensiva®SC50, at levels of typically 1-15 percent. Compositions disclosed in United States Patent No. 5,885,562 may further comprise a short chain monohydric alcohol such as ethanol at a level of between 20 and 80 percent. Formulations useful as deodorants, however, would differ from those used as skin sanitizers in that skin sanitizers would optimally exhibit rapid broad spectrum activity against bacteria, fungi, and viruses, not merely gram positive odor causing bacteria.

[0015] United States Patent No. 5,516,510 by Beilfuss et al., issued May 14, 1996, discloses deodorant compositions which comprise glycerin monoalkyl ethers such as octoxyglycerin (referred to therein as 2-ethyl hexyl glycerin ether, and as being the most preferred among these compounds). The deodorant compositions of United States Patent No. 5,516,510 may be formulated in aqueous and/or alcoholic solutions and may further comprise additional antimicrobial compounds, including triclosan, chlorhexidine salts, alexidine salts, and

phenoxyethanol, among others. Specific concentration ranges for triclosan and the biguanides are not provided.

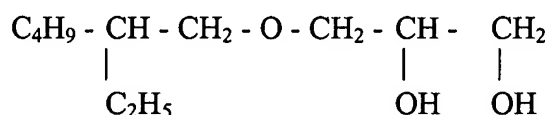
3. SUMMARY OF THE INVENTION

[0016] The present invention relates to antimicrobial compositions comprising synergistic combinations of octoxyglycerin and at least one other antimicrobial agent in formulations which are more effective than prior art compositions without causing increased irritation to the skin of the average user. In certain embodiments, skin irritation may be minimized by low concentrations of antimicrobials and/or the presence of soothing compounds such as zinc. Preferred embodiments of the invention comprise combinations of octoxyglycerin, a quaternary ammonium compound, and at least one other antimicrobial agent. Without being bound to any particular theory, it is hypothesized that the unexpected antimicrobial effectiveness of combinations of octoxyglycerin may result from an enhancement of the permeability of microbes to antimicrobials caused by octoxyglycerin.

4. DETAILED DESCRIPTION OF THE INVENTION

[0017] The present invention relates to combinations of octoxyglycerin with at least one, and preferably at least two, antimicrobial agents. In preferred embodiments of the invention such compositions comprise octoxyglycerin and a quaternary ammonium compound.

[0018] Octoxyglycerin, as used herein, is also known as glycerol 1-(2-ethylhexyl) ether and is sold under the trade name Sensiva® SC 50 ("Sensiva®") by Schulke & Mayr (Rockaway, New Jersey). Octoxyglycerin has the following chemical structure:



which has the empirical formula $C_{11}H_{24}O_3$. The CAS No. of octoxyglycerin is 70445-33-9.

Octoxyglycerin has a relative molecular weight of 204.31 g/mol. Sensiva® SC 50 is sold as a clear, almost colorless liquid, having a refractive index of approximately 1.451, a density at 20°C of approximately 0.95 g/ml, a boiling point of >285°C, a flash point of 152°C, a water solubility at 22°C of approximately 1.8 g/l and virtually complete solubility in fat. In addition to having antimicrobial activity, it acts as a mild humectant and skin emollient. The present invention provides for compositions comprising octoxyglycerin at between 1 and 5 percent, and preferably 1-3 percent. It should be noted that all ranges recited herein are inclusive of their limiting values. Sensiva SC50 is essentially pure octoxyglycerin.

[0019] Antimicrobial agents which may be used in addition to octoxyglycerin according to the invention include biguanides and phenols. Biguanides may be used in concentrations between about 0.05 and 4 percent and preferably between about 0.05 and 2 percent. Examples of suitable biguanides include polyhexamethylene biguanide (PHMB) at concentrations between about 0.3 and 1 percent, alexidine at concentrations between about 0.5 and 2 percent, and chlorhexidine compounds at concentrations between about 0.5 and 4 percent and preferably between about 0.05 and 1 percent. A chlorhexidine compound, as that term is used herein, includes chlorhexidine free base as well as chlorhexidine salts, including, but not limited to, chlorhexidine diacetate (also known as "chlorhexidine acetate"), chlorhexidine digluconate (also known as "chlorhexidine gluconate"), chlorhexidine palmitate, chlorhexidine diphosphanilate, chlorhexidine dihydrochloride, chlorhexidine dichloride, chlorhexidine dihydroiodide, chlorhexidine diperchlorate, chlorhexidine dinitrate, chlorhexidine sulfate, chlorhexidine sulfite, chlorhexidine thiosulfate, chlorhexidine di-acid phosphate, chlorhexidine difluorophosphate, chlorhexidine diformate, chlorhexidine dipropionate, chlorhexidine di-iodobutyrate,

chlorhexidine di-n-valerate, chlorhexidine dicaproate, chlorhexidine malonate, chlorhexidine succinate, chlorhexidine malate, chlorhexidine tartrate, chlorhexidine dimonoglycolate, chlorhexidine monodiglycolate, chlorhexidine dilactate, chlorhexidine di-alpha-hydroxyisobutyrate, chlorhexidine diglucoheptonate, chlorhexidine di-isothionate, chlorhexidine dibenzoate, chlorhexidine dicinnamate, chlorhexidine dimandelate, chlorhexidine di-isophthalate, chlorhexidine di-2-hydroxynapthoate, and chlorhexidine embonate. Most preferably, the chlorhexidine compound is chlorhexidine digluconate a concentration between 0.05 and 4 percent.

[0020] Phenols (phenol derivatives) which may be used according to the invention include, but are not limited to, 2-hydroxyphenol compounds such as triclosan (2,4,4'-trichloro-2'-hydroxydiphenyl ether, also available as IRGASAN DP300 from Ciba Specialty Chemicals Corp, Greensboro, NC) and 2,2'-dihydroxy-5,5'-dibromo-diphenyl ether; p-nitrophenol, picric acid, xlenol, phenoxyethanol, chlorinated phenols such as parachlorometaxlenol, p-chloro-o-benzylphenol and-dichlorophenol, cresols such as p-chloro-m-cresol, pyrocatechol, resorcinol, 4-n-hexylresorcinol, pryogallol, phloroglucin, carvacrol, thymol, p-chlorothymol, o-phenylphenol, o-benzylphenol, phenol, 4-ethylphenol, 4-phenolsulfonic acids, hexachlorophene, tetrachlorophene, dichlorophene, 2,3-dihydroxy-5,5'-dichlorophenyl sulfide, 2,2'-dihydroxy-3,3',5,5'-tetrachlorodiphenyl sulfide, 2,2'-dihydroxy-3,3',5,5',6,6'-hexachlorodiphenyl sulfide and 3,3'-dibromo-5,5'-dichloro-2,2'-dihydroxydiphenylamine. Preferred is triclosan at a concentration of between about 0.1 and 2 percent and most preferably between about 0.3 and 1 percent. Other phenols may be comprised at concentrations of between about 0.3 and 2 percent, but preferably at concentrations equivalent in potency against *S. aureus* as between 0.3 and 1 percent triclosan.

[0021] Additional antimicrobial agents which may be incorporated into compositions of the invention include antifungal agents such as miconazole (preferably at a concentration of 1 - 2 percent), polymixin (preferably at a concentration of 0.3 - 1 percent), neomycin (preferably at a concentration of 0.1 - 0.5 percent), iodine compounds such as povidone iodine (preferably at a concentration of 1 - 10 percent), minocycline (preferably at a concentration of 0.3 - 1.0 percent), and metal salts such as silver sulfadiazine (preferably at a concentration of 1 - 2 percent).

[0022] Preferred non-limiting embodiments of the invention comprise octoxyglycerin together with a quaternary ammonium compound, such as, but not limited to, benzalkonium chloride ("BZK", which is particularly preferred), benzethonium chloride, other benzalkonium or benzethonium halides, including, but not limited to, benzalkonium or benzethonium bromide or fluoride, cetyl pyridinium chloride, dequalinium chloride, N-myristyl-N-methyl-morpholinium methyl sulfate, poly[N-[3-(dimethylammonio)propyl]-N'-[3-(ethyleneoxyethyl)dimethylammonio]propyl]urea dichloride], alpha-4-[1-tris(2-hydroxyethyl)ammonium chloride-2-butenyl]-omega-tris(2-hydroxyethyl)ammonium chloride, poly[oxyethylene (dimethyliminio)ethylene (dimethyliminio)-ethylene dichloride]. The concentrations of quaternary ammonium compound may be between about 0.01 and 0.3 percent; preferably the quaternary ammonium compound is benzalkonium chloride at a concentration between 0.05 and 0.2 percent, more preferably between 0.1 and 0.15 percent.

[0023] In certain non-limiting embodiments, compositions of the invention may further comprise one or more alcohol. Alcohols which may be used according to the invention include aliphatic alcohols, including, but not limited, most preferred ethanol or isopropyl alcohol, but also n-propyl alcohol, and mixtures thereof, at concentrations between about 20 and 85 percent and preferably 40 to 70 percent. Suitable alcohols also include fatty alcohols, such as cetyl

alcohol, myristyl alcohol, stearyl alcohol, octyl alcohol, decyl alcohol, lauryl alcohol, and combinations thereof, at concentrations between about 0.5 and 5 percent. The present invention further provides for compositions comprising, as at least one alcoholic component, hexanol at a concentration of between three and ten percent and preferably about 5 percent.

[0024] The formulations of the invention may further comprise one or more of the following:

A zinc-containing compound such as a zinc salt, including but not limited to zinc gluconate, zinc oxide, zinc stearate, zinc salicylate, zinc carbonate, zinc oleate, zinc acetate, zinc peroxide, zinc phosphate, and zinc undecylenate. Zinc compounds are known to have anti-irritant activity (see, for example, United States Patent No. 5,965,610 by Modak et al. and U.S. Patent No. 5,985,918 by Modak et al., incorporated by reference herein). Preferred zinc compounds for use according to the invention are, for a disinfecting alcohol gel, zinc gluconate and zinc oxide, at concentrations between 0.1 and 1 percent, and preferably 0.8 percent zinc gluconate and 0.2 percent zinc oxide; for an antiseptic aqueous formulation, zinc gluconate and zinc stearate, at concentrations between 0.2 and 7 percent, and preferably 2.4 percent zinc gluconate and 3.8 percent zinc stearate.

An emollient, which may be, for example, an organic, a hydrocarbon-based or a fatty-ester based emollient. Suitable hydrocarbon-based emollients include petrolatum and mineral oils. Suitable fatty ester based emollients include methyl, isopropyl and butyl esters of fatty acids such as isopropyl palmitate, isopropyl myristate, isopropyl isostearate, isostearyl isostearate, diisopropyl sebacate, and propylene dipelargonate, 2-ethylhexyl isononoate, 2-ethylhexyl stearate, C₁₂ - C₁₆ fatty alcohol lactates such as cetyl lactate and lauryl lactate, isopropyl lanolate, 2-ethylhexyl salicylate, cetyl myristate, oleyl myristate, oleyl stearate, oleyl

oleate, hexyl laurate, and isohexyl laurate. Additional useful emollients include lanolin, olive oil, cocoa butter, and shea butter.

A humectant, such as, for example, glycerine, 1-2-propylene glycol, dipropylene glycol, polyethylene glycol, 1,3-butylene glycol, or 1,2,6-hexanetriol.

A thickening and/or gelling agent, such as, for example, an addition polymer of acrylic acid, a resin such as Carbopol® ETD™ 2020, guar gum, acacia, acrylates/stearate-20 methacrylate copolymer, agar, algin, alginic acid, ammonium acrylate co-polymers, ammonium alginate, ammonium chloride, ammonium sulfate, amylopectin, attapulgate, bentonite, C9-15 alcohols, calcium acetate, calcium alginate, calcium carrageenan, calcium chloride, caprylic alcohol, carbomer 910, carbomer 934, carbomer 934P, carbomer 940, carbomer 941, carboxymethyl hydroxyethyl cellulose, carboxymethyl hydroxypropyl guar, carrageenan, cellulose, cellulose gum, cetearyl alcohol, cetyl alcohol, corn starch, damar, dextrin, dibenzylidene sorbitol, ethylene dihydrogenated tallowamide, ethylene diolamide, ethylene distearamide, gelatin, guar gum, guar hydroxypropyltrimonium chloride, hectorite, hyaluronic acid, hydrated silica, hydroxybutyl methylcellulose, hydroxyethylcellulose, hydroxyethyl ethylcellulose, hydroxyethyl stearamide-MIPA, hydroxypropylcellulose, hydroxypropyl guar, hydroxypropyl methylcellulose, isocetyl alcohol, isostearyl alcohol, karaya gum, kelp, lauryl alcohol, locust bean gum, magnesium aluminum silicate, magnesium silicate, magnesium trisilicate, methoxy PEG-22/dodecyl glycol copolymer, methylcellulose, microcrystalline cellulose, montmorillonite, myristyl alcohol, oat flour, oleyl alcohol, palm kernel alcohol, pectin, PEG-2M, PEG-5M, polyacrylic acid, polyvinyl alcohol, potassium alginate, potassium aluminum polyacrylate, potassium carrageenan, potassium chloride, potassium sulfate, potato starch, propylene glycol alginate, sodium acrylate/vinyl alcohol copolymer, sodium carboxymethyl dextran, sodium

carrageenan, sodium cellulose sulfate, sodium chloride, sodium polymethacrylate, sodium silicoaluminate, sodium sulfate, stearylalkonium bentonite, stearylalkonium hectorite, stearyl alcohol, tallow alcohol, TEA-hydrochloride, tragacanth gum, tridecyl alcohol, tromethamine magnesium aluminum silicate, wheat flour, wheat starch, xanthan gum, abietyl alcohol, acrylinoleic acid, aluminum behenate, aluminum caprylate, aluminum dilinoleate, aluminum salts, such as distearate, and aluminum isostearates, beeswax, behenamide, behenyl alcohol, butadiene/acrylonitrile copolymer, C29-70 acid, calcium behenate, calcium stearate, candelilla wax, carnauba, ceresin, cholesterol, cholesterol hydroxystearate, coconut alcohol, copal, diglyceryl stearate malate, dihydroabietyl alcohol, dimethyl lauramine oleate, dodecanoic acid/cetearyl alcohol/glycol copolymer, erucamide, ethylcellulose, glyceryl triacetate hydroxystearate, glyceryl tri-acetyl ricinolate, glycol dibehenate, glycol di-octanoate, glycol distearate, hexanediol distearate, hydrogenated C6-14 olefin polymers, hydrogenated castor oil, hydrogenated cottonseed oil, hydrogenated lard, hydrogenated menhaden oil, hydrogenated palm kernel glycerides, hydrogenated palm kernel oil, hydrogenated palm oil, hydrogenated polyisobutene, hydrogenated soybean oil, hydrogenated tallow amide, hydrogenated tallow glyceride, hydrogenated vegetable glyceride, hydrogenated vegetable oil, Japan wax, jojoba wax, lanolin alcohol, shea butter, lauramide, methyl dehydroabietate, methyl hydrogenated rosin, methyl rosin, methylstyrene/vinyltoluene copolymer, microcrystalline wax, montan acid wax, montan wax, myristyleicosanol, myristyloctadecanol, octadecene/maleic anhydride copolymer, octyldodecyl stearyl stearate, oleamide, oleostearine, ouricury wax, oxidized polyethylene, ozokerite, paraffin, pentaerythrityl hydrogenated rosin, pentaerythrityl tetraoctanoate, pentaerythrityl rosin, pentaerythrityl tetraabietate, pentaerythrityl tetrabehenate, pentaerythrityl tetraoleate, pentaerythrityl tetrastearate, phthalic anhydride/glycerine/glycidyl

decanoate copolymer, phthalic/trimellitic/glycols copolymer, polybutene, polybutylene terephthalate, polydipentene, polyethylene, polyisobutene, polyisoprene, polyvinyl butyral, polyvinyl laurate, propylene glycol dicaprylate, propylene glycol dicocoate, propylene glycol diisononanoate, propylene glycol dilaurate, propylene glycol dipelargonate, propylene glycol distearate, propylene glycol diundecanoate, PVP/eiconsene copolymer, PVP/hexadecene copolymer, rice bran wax, stearylkonium bentonite, stearylkonium hectorite, stearamide, stearamide DEA-distearate, stearamide DIBA-stearate, stearamide MEA-stearate, stearone, stearyl alcohol, stearyl erucamide, stearyl stearate, stearyl stearyl stearate, synthetic beeswax, synthetic wax, trihydroxystearin, triisononanoate, triisostearin, tri-isostearyl trilinoleate, trilaurin, trilinoleic acid, trilinolein, trimyristin, triolein, tripalmitin, tristearin, zinc laurate, zinc myristate, zinc neodecanoate, zinc rosinate, and mixtures thereof.

A neutralizing agent, which may be included, for example, to neutralize carboxyl groups present in one or more other component, such as carboxyl groups in a thickening agent. Suitable neutralizing agents include diisopropylamine and triethanolamine.

A surfactant, which may be an anionic surfactant, a cationic surfactant, an ampholytic surfactant, or a nonionic surfactant, such as, for example, nonionic surfactants such as polyethoxylates, fatty alcohols (e.g., ceteth-20 (a cetyl ether of polyethylene oxide having an average of about 20 ethylene oxide units) and other "BRIJ"® nonionic surfactants available from ICI Americas, Inc. (Wilmington, DE)), cocamidopropyl betaine, alkyl phenols, fatty acid esters of sorbitol, sorbitan, or polyoxyethylene sorbitan. Suitable anionic surfactants include ammonium lauryl sulfate and lauryl ether sulfosuccinate. A preferred surfactant is lauroyl ethylenediamine triacetic acid sodium salt at a concentration between about 0.5 - 2.0%. Suitable concentrations of surfactant are between about 0.05 and 2 percent.

Water used in the formulations is preferably deionized water having a neutral pH.

Additional additives, including but not limited to a silicone fluid (such as dimethicone or cyclomethicone), dyes, fragrances, etc. Examples of additional additives include but are not limited to: pH adjusters, including basic pH adjusters such as ammonia, mono-, di- and tri- alkyl amines, mono-, di- and tri-alkanolamines, alkali metal and alkaline earth metal hydroxides (*e.g.*, ammonia, sodium hydroxide, potassium hydroxide, lithium hydroxide, monoethanolamine, triethylamine, isopropylamine, diethanolamine and triethanolamine); acid pH adjusters such as mineral acids and polycarboxylic acids (*e.g.*, hydrochloric acid, nitric acid, phosphoric acid, sulfuric acid, citric acid, glycolic acid, and lactic acid); vitamins such as vitamin A, vitamin E and vitamin C; polyamino acids and salts, such as ethylenediamine tetraacidic acid (EDTA), preservatives such as DMDM hydantoin, and sunscreens such as aminobenzoic acid, arobenzone, cinoxate, dioxybenzone, homosalate, menthyl anthranilate, octocrylene, octyl methoxycinnamate, octyl salicylate, oxybenzoate, padimate O, phenylbenzimidazole, sulfonic acid, sulisobenzene, titanium dioxide, trolamine salicylate and zinc oxide.

[0025] Specific, non-limiting embodiments of the invention include the following compositions, which may further comprise additional ingredients which do not substantially effect the antimicrobial properties of the composition. For the following formulations, the water indicated was added last to the other ingredients to bring the total volume to 100 percent.

1. An antiseptic alcohol gel comprising:

zinc gluconate	0.8 percent
zinc oxide	0.2 percent
ethyl alcohol	65.0 percent (volume/volume)
hydroxy methyl propyl cellulose (K100M)	0.3 percent

Ucare JR 400 (polyquaternium-10) (Amerchol Corp.)	0.15 percent
Incroquat Behenyl TMS (Croda, Inc.)	1.0 percent
Polawax A-31 (Croda, Inc.)	1.0 percent
stearyl alcohol - Crodacol(S70) (Croda, Inc.)	1.0 percent
Cremerol HMG (Amerchol Corp.)	1.0 percent
dimethicone	0.5 percent (volume/volume)
Germall plus (ISP Sutton Laboratories)	0.25 percent
propylene glycol	1.5 percent (volume/volume)
glycerine	1.0 percent (volume/volume)
water	23.13 percent (volume/volume)
chlorhexidine digluconate	0.05 percent
phenoxyethanol	1.0 percent
BZK	0.12 percent
Sensiva SC50	2 percent (volume/volume)

where the gel may be applied to and rubbed over the skin to achieve its antimicrobial effect.

2. An antiseptic alcohol gel comprising:

water	31.32 percent (volume/volume)
Ucare (Amerchol Corp.)	0.08 percent

hydroxypropylmethylcellulose (K-100) (Dow Corning)	0.15 percent
Polyox WSR 301 (polyethyleneoxide) (Dow Corning)	0.03 percent
Incroquat (Croda, Inc.)	0.4 percent
Polawax A-31 (Croda, Inc.)	0.4 percent
polyethylene glycol	0.25 percent
ethanol	63.5 percent (volume/volume)
Glucam E-20 (Amerchol Corp.)	0.4 percent
Silicone 225 (Dow Corning)	0.1 percent (volume/volume)
Sensiva SC50	2.0 percent (volume/volume)
phenoxyethanol	1.0 percent
chlorhexidine digluconate	0.05 percent
BZK	0.12 percent
Germall Plus (Sutton Laboratories)	0.2 percent

3. An antiseptic aqueous formulation comprising:

zinc gluconate	2.4 percent
zinc stearate	3.8 percent
hydroxy methyl propyl cellulose (K100M)	0.5 percent
Kytamer PC (Chitisan) (Amerchol Corp.)	0.15 percent

Ucare JR 400 (Amerchol Corp.)	0.1 percent
Incroquat behenyl TMS (Croda, Inc.)	1.0 percent
Crodamol NM (Croda, Inc.)	1.6 percent
Acetulan (Amerchol Corp.)	2.0 percent
Cremerol HMG (Amerchol Corp.)	1.0 percent
stearyl alcohol	2.0 percent
allantoin	0.25 percent
Germall Plus (ISP Sutton Laboratories)	0.3 percent
dimethicone	1.0 percent (volume/volume)
water	81.48 percent (volume/volume)
PHMB	0.3 percent
phenoxyethanol	1.0 percent
BZK	0.12 percent
Sensiva SC50	2 percent (volume/volume)

4. An antimicrobial scrub gel comprising:

water	30.5 percent
Ucare (Amerchol Corp.)	0.1 percent
hydroxy propyl methyl cellulose (K100) (Dow Corning)	0.2 percent

Polyox WSR 301 (polyethyleneoxide) (Dow Corning)	0.1 percent
Incroquat (Croda, Inc.)	0.4 percent
Polawax A-31 (Croda, Inc.)	0.4 percent
propylene glycol	1.0 percent
ethanol	63.5 percent (volume/volume)
Glucam E-20 (Amerchol Corp.)	0.4 percent
Masil SF 19 CG surfactant	1.0 percent
phenoxyethanol	1.0 percent
Sensiva SC50	1.0 percent (volume/volume)
chlorhexidine digluconate	0.05 percent
BZK	0.12 percent
Germall Plus (Sutton Laboratories)	0.2 percent

5. An antimicrobial scrub gel, for example for pre-operative skin disinfection,
comprising:

ethanol	35 percent (volume/volume)
isopropanol	35 percent (volume/volume)
zinc gluconate	0.5 percent
zinc oxide	0.2 percent
hydroxy methyl propyl cellulose (K100M)	0.3 percent
Germall Plus (ISP Sutton Laboratories)	0.25 percent

hexanol	5.0 percent (volume/volume)
PXE	1.0 percent
Sensiva	1.5 percent (volume/volume)
chlorhexidine digluconate	0.05 percent

with water added to 100 percent (approximately 21.2 milliliters/100 ml solution).

6. Another antimicrobial scrub gel, for example for pre-operative skin disinfection, comprising:

water	23.28 percent (volume/volume)
Polyox WSR 205	0.2 percent
U-care JR 400	0.2 percent
ethanol (95%)	65 percent (volume/volume)
propylene glycol	3 percent
Sensiva SC50	2 percent (volume/volume)
BZK	0.12 percent
phenoxyethanol	1.0 percent
povidone iodine	5.0 percent
Germall Plus	0.2 percent

7. An antimicrobial soap comprising:

water	51.2 percent (volume/volume)
Ucare (Amerchol Corp.)	0.1 percent
hydroxy propyl methyl cellulose (K-100) (Dow Corning)	0.2 percent
Polyox WSR 301 (polyethyleneoxide)	0.03 percent

ethanol	40 percent (volume/volume)
Pluronic F-87 (BASF)	2.0 percent
Masil SF 19 CG surfactant	1.0 percent
Cocamidopropyl betaine (Witco Corp.)	2.0 percent
propylene glycol	1.0 percent
phenoxyethanol	1.0 percent
chlorhexidine digluconate	0.05 percent
BZK	0.12 percent
Sensiva SC50	0.5 percent (volume/volume)
Germall Plus (Sutton Laboratories)	0.2 percent

8. An antifungal cream comprising miconazole (1-2 percent), chlorhexidine digluconate (0.05 - 0.2 percent), and Sensiva SC50(1-3 percent) in a hydrophilic cream base.

9. A topical antiseptic ointment for wound care comprising polymixin (0.3 - 1%), neomycin (0.1 - 0.5 percent), chlorhexidine digluconate (0.05 - 0.2 percent), and Sensiva SC50(1 - 3 percent) in a hydrophilic base.

10. A topical antiseptic ointment for burn wound care comprising silver sulfadiazine (1 - 2 percent), chlorhexidine digluconate (0.05 - 0.2 percent) and Sensiva SC50 (1 - 3 percent) in a hydrophilic base.

5. EXAMPLES

EXAMPLE 1 : SENSIVA + BZK

[0026] Sensiva SC50 and/or benzalkonium chloride ("BZK") were added, in various

concentrations, to the following alcohol gel base:

ethyl alcohol	65 percent (volume/volume)
hydroxy methyl propyl cellulose (K100M)	0.3 percent
hydroxy propyl cellulose (HF)	0.1 percent (volume/volume)
Glucam P20	1.0 percent (volume/volume)
Glucam P20 distearate	1.5 percent (volume/volume)
Ucare JR 400 (polyquaternium-10)	0.15 percent
silicone (DC 1403)	1.5 percent (volume/volume)
Germall Plus	0.25 percent

to which water was added, after the incorporation of other additives, to bring the total volume to 100 percent (typically requiring approximately 20-30 percent (volume/volume)).

The amount of Sensiva, throughout the example section, is a volume/volume percentage.

[0027] Antimicrobial activity was evaluated using the following assay. 1 milliliter of 10^8 colony forming units ("cfus") of test organism per milliliter was added to 1 milliliter of bovine adult serum in a sterile culture tube and mixed. 1 milliliter of the test gel was added to each tube, and was vortexed to mix. After 15 seconds, three 0.5 ml aliquots were removed and further diluted 1:1000 with LTSB (lecithin-containing trypticase soy broth) drug inactivating medium, and, of the resulting liquid, 0.5 milliliters were plated on each trypticase soy agar ("TSA") plate. The resulting plates were incubated at 37° C for 24 hours and the colony count per tube was determined.

[0028] The foregoing method was used to determine the antimicrobial activities of formulations of the above alcohol gel base comprising either Sensiva SC50, BZK or combinations of Sensiva SC50 and BZK. The results for Sensiva SC50 used alone are shown in

Table 1, and the results for Sensiva SC50, BZK and Sensiva SC50/BZK combinations are shown in Table 2.

TABLE 1

% Sensiva	0	0.5	1.0	2.0	3.0	5.0
<i>S. aureus</i> (cfu/tube)	1×10^8	1×10^7	4×10^7	3×10^6	1×10^6	1×10^6
fold-reduction*	-	2.5	9	33	100	100

*relative to control

TABLE 2

% Sensiva	0	1.0	2.0	0	0	0	1.0	1.0	2.0	2.0
% BZK	0	0	0	0.12	0.19	0.5	0.12	0.19	0.12	0.19
<i>S. aureus</i> (cfu/tube)	1×10^8	4×10^7	3×10^6	1.6×10^7	2×10^7	3.7×10^6	8×10^5	2×10^4	8×10^3	3.0×10^3
Log 10 cfu reduction relative to control	-	1	1.5	0.8	0.7	1.4	2.1	3.7	4.1	4.5
Increase in log 10 beyond additive effect	NA	NA	NA	NA	NA	NA	0.3	2	1.8	2.3
fold reduction relative to control	-	10	33	6.25	5	27	125	5×10^3	1.25×10^4	3.3×10^4

[0029] Tables 1 and 2 show that no significant antimicrobial activity against *S. aureus* was obtained with 2-5 percent Sensiva; the antimicrobial activity was not significantly different between 2, 3 and 5 percent of Sensiva. Similarly, 0.12 and 0.19 percent BZK exhibited minimal or no antimicrobial activity (Table 2). However, combinations of 1-2 percent Sensiva SC50 and 0.12-0.19 percent BZK showed 5000-33000 fold reduction in colony forming units compared to control values (Table 2).

EXAMPLE 2 : SENSIVA + CHLORHEXIDINE DIGLUCONATE

[0030] Assays using the same gel base and protocol as set forth in Example 1 to test activities of Sensiva, chlorhexidine digluconate ("CHG"), and combinations thereof gave the following results, shown in Table 3.

TABLE 3

% Sensiva	0	0	0	0	1.0	1.0	1.0	2.0	2.0	2.0
% CHG	0	0.05	0.25	0.5	0.05	0.25	0.5	0.05	0.25	0.5
<i>S. aureus</i> (cfu/tube)	1×10^8	1.1×10^7	8×10^6	4.2×10^6	1.2×10^5	6×10^4	8×10^3	8×10^3	5×10^3	1×10^3
Log 10 cfu reduction relative to control	-	1.0	1.1	1.4	2.9	3.1	4.1	4.1	4.3	4.3
Increase in log 10 beyond additive effect	NA	NA	NA	NA	0.4	1.1	1.7	1.6	1.7	2.4
fold reduction relative to control	-	9	12.5	23.8	833	1666	12500	12500	20000	1×10^5

Thus, Sensiva SC50 (1-2 percent) and CHG (0.05-0.5 percent) used individually showed 9-35 fold reduction in colony counts as compared to control, whereas a combination of 1-2 percent Sensiva with 0.05-0.5 percent CHG showed 800-100,000 fold reduction. Thus, the combination of Sensiva and CHG appears to be synergistic. When benzalkonium chloride was added to formulation, the antimicrobial activity was improved still further, as shown in the following example section.

EXAMPLE 3 : SENSIVA + CHLORHEXIDINE DIGLUCONATE + BZK

[0031] Assays using the same gel base and protocol as set forth in Example 1 to test activities of combinations of Sensiva, chlorhexidine digluconate ("CHG") and BZK gave the following results, shown in Table 4.

TABLE 4

% Sensiva	0	0	1.0	2.0
% BZK	0	0.12	0.12	0.12
% CHG	0	0.05	0.05	0.05
Growth (cfu/ml)	1×10^8	1.2×10^7	4×10^4	0
Log 10 cfu reduction relative to control	0	1.0	4.0	8.0
Increase in log 10 beyond additive effect	NA	NA	2.1	5.1
fold reduction relative to control	-	8.3	2500	10^8

NA = not applicable

EXAMPLE 4: COMBINATIONS OF SENSIVA AND OTHER ANTIMICROBIALS

[0032] Since Sensiva does not exhibit potent microbicidal activity even at concentrations of between 3 and 5 percent, it is surprising that this compound exhibits synergism with chlorhexidine digluconate and BZK. Octoxyglycerin (Sensiva) has been reported to have the property of deeper penetration into the upper layers of the epidermis. Without being bound by any particular theory, the mechanism of synergistic action may be explained as follows. When a bacterium is exposed to Sensiva and a second antimicrobial agent, Sensiva may penetrate through the bacterial cell wall and thereby compromise the bacterial transport system. This may result in increased uptake of the second antimicrobial agent. This mechanism would indicate

that Sensiva would promote the antimicrobial effects of a diverse array of compounds, including quaternary ammonium compounds, biguanides, chlorinated phenols, metal salts, antifungal azoles, etc..

[0033] Accordingly, the antimicrobial activity of various combinations of Sensiva and other antimicrobials was tested, using concentrations that fall within the recommended usage range for topical formulations. The following agents were tested. Benzalkonium chloride (BZK) and benzethonium chloride (BZT) were tested as representative of the class of quaternary ammonium compounds. Chlorhexidine digluconate (CHG) and polyhexamethylene biguanide (PHMB) were tested as representative of the class of biguanides. Parachlorometaxylenol (PCMX) and triclosan (TC) were tested as representative of the class of chlorinated phenols. Povidone iodine (PVI) was tested as representative of the class of iodine compounds. Silver sulfadiazine (AgSD) was tested as representative of the class of metal salts. Neomycin and miconazole were tested as representative of the class of antibiotics. The alcohol gel base and protocol set forth in Example 1 were used to produce the data set forth in Table 5.

[0034] Similar protocols were then used to test the antibacterial activity of Sensiva combined with chlorhexidine digluconate and another antimicrobial agent. The results are shown in Table 6.

TABLE 5

% Antimicrobial	% Sensiva	Growth (CFU/ml)	fold reduction*
0 Control	0	1×10^8	-
0	2.0	3×10^6	33
BZK			
0.12	0	1.6×10^7	6.25
0.12	2.0	8.0×10^3	12500
BZT			
0.12	0	1.0×10^7	10
0.12	2.0	5.0×10^3	20,000
CHG			
0.05	0	1.1×10^7	9
0.05	2.0	8.0×10^3	12,500
PHMB			
0.3	0	3.0×10^6	33
0.3	2.0	4.0×10^3	25,000
TC			
0.3	0	1.0×10^8	0
0.3	2.0	2.2×10^5	450
PCMX			
0.3	0	1.0×10^8	0
0.3	2.0	6.2×10^4	1612
AgSD			
1.0	0	1.0×10^8	0
1.0	2.0	3.0×10^5	330
PVI			
1.0	0	2.0×10^7	5
1.0	2.0	3.0×10^4	3,333
Neomycin			
0.3	0	2.3×10^7	4.3
0.3	2.0	1.0×10^3	100,000
Miconazole			
1.0	0	1.0×10^8	0
1.0	2.0	6.0×10^4	1666

*relative to control

TABLE 6

% Antimicrobial	% Sensiva	%CHG	Growth (CFU/ML)	Fold Reduction Compared to Control
0	0	0	1.0×10^8	-
0	2.0	0	3.0×10^6	33
0	2.0	0.05	8.0×10^3	12,500
BZK				
0.12	0	0.05	1.2×10^7	8.3
0.12	2.0	0.05	0	10^8
TC				
0.3	0	0.05	9.0×10^6	11.1
0.3	2.0	0.05	0	10^8
PCMX				
0.3	0	0.05	7.0×10^6	14.2
0.3	2.0	0.05	0	10^8
AgSD				
1.0	0	0.05	1.0×10^7	10
1.0	2.0	0.05	0	10^8
PVI				
1.0	0	0.05	1.0×10^7	10
1.0	2.0	0.05	0	10^8
Neomycin				
0.3	0	0.05	1.0×10^6	100
0.3	2.0	0.05	0	10^8

[0035] The data shown in Table 5 indicate that Sensiva, at a concentration of 2.0 percent, produced a 33-fold reduction in bacterial colony formation, and the antibacterial activity of the other antimicrobials tested, used alone, was less than or equal to 33-fold. Combination of these antimicrobials with Sensiva greatly resulted in an antibacterial activity greater than what would have been expected, based on the inhibitory activity of either agent used separately. The extent of this enhancement varied among antimicrobials; for example, the activity of quaternary ammonium compounds, used in combination with Sensiva, was observed to be 12,500 and 20,000-fold greater than control. The biguanides chlorhexidine digluconate and parahexamethylenebiguanide, in combination with Sensiva, produced an antimicrobial activity

12,500 and 25,000-fold greater, respectively, than control. Neomycin, in combination with Sensiva, exhibited an antimicrobial activity 100,000 greater than control. Thus, Sensiva has been demonstrated to enhance the antimicrobial effects of a wide variety of agents. The data shown in Table 6 further show that combinations of Sensiva and chlorhexidine digluconate with various antimicrobials exhibit a further enhancement in activity.

EXAMPLE 5 : ADDITIONAL DATA

[0036] Assays using the same gel base and protocol as set forth in Example 1 to test activities of combinations of Sensiva and other antimicrobials gave the following results, shown in Table 7.

TABLE 7

Agent(s)	Concentrations	Growth (cfu/tube)
control (without gel base)	-	$2.5 - 4.2 \times 10^8$
Sensiva	0.5	4.0×10^7
Sensiva	1.0	1.0×10^7
BZK	0.019	8.0×10^7
BZK + Sensiva	0.019 1.0	2.0×10^7
BZK + Sensiva	0.019 2.0	1.2×10^7
BZK	0.12	1.6×10^7
BZK + Sensiva	0.12 0.5	1.4×10^7
BZK + Sensiva	0.12 1.0	8.0×10^5
CHG	0.05	1.1×10^7
CHG + Sensiva	0.05 0.5	6.3×10^6
CHG + Sensiva	0.05 1.0	1.2×10^5
PCMX	0.15	3.5×10^8
PCMX + Sensiva	0.15 2.0	4.1×10^5
TC + BZK	0.3 0.12	1.0×10^7
TC + BZK + Sensiva	0.3 0.12 2.0	4.0×10^3
PCMX + BZK	0.3 0.12	2.0×10^6
PCMX + BZK + Sensiva	0.3 0.12 2.0	1.0×10^3
Miconazole + CHG	1.0 0.05	1.0×10^7
Miconazole + CHG + Sensiva	1.0 0.05 2.0	1.0×10^3
PVI + CHG	1.0 0.05	1.0×10^7
PVI + CHG + Sensiva	1.0 0.05 2.0	0

EXAMPLE 6: COMBINATIONS OF SENSIVA, BZK, AND OTHER AGENTS

[0037] Again using the alcohol gel base and protocol described in Example 1, various combinations of Sensiva, the quaternary ammonium compound BZK, and other antimicrobials produced the results shown in Table 8.

TABLE 8

Agent(s)	Concentration (%)	Growth (cfu/tube)
Control (no gel base)	-	2.0×10^8
Control (gel base)	-	1.2×10^8
PXE	1.0	1.0×10^8
PXE + Sensiva	1.0 1.0	2.0×10^7
PXE + Sensiva	1.0 2.0	3.3×10^5
BZK + CHG + Sensiva	0.12 0.05 1.0	4.0×10^4
BZK + CHG + Sensiva	0.12 0.05 2.0	0
BZK + CHG + Sensiva + PXE	0.12 0.05 1.0 1.0	0
BZK + PHMB + Sensiva	0.12 0.3 1.0	8.0×10^3
BZK + PHMB + Sensiva + PXE	0.12 0.3 1.0 1.0	0

[0038] The above data demonstrates that the addition of the phenol derivative, phenoxyethanol, enhanced the antimicrobial activity of several combinations of Sensiva and other antimicrobials.

EXAMPLE 7: SUSTAINED ACTIVITY OF ANTIMICROBIAL PREPARATIONS

[0039] Natural leather was cut into 2 x 2 cm pieces, washed, and sterilized. For each test group 4 pieces were used. Equal amounts (0.25 ml) of various test formulations were applied uniformly on the surface of each piece, and then allowed to dry for 3 hours. Then 10 microliters of a *Staphylococcus aureus* culture (10^7 CFU/ml) was spread uniformly on the surface of the treated leather patches. After 1 minute, the inoculated side of the leather was rinsed with 10 ml of drug inactivating medium (LTSB), of which a 0.5 ml aliquot was plated on the surface of a D/E (drug inactivating) plate. Plates prepared in this manner were incubated for 24 hours at 37°C and bacterial colonies were counted. The results, which demonstrate sustained antimicrobial activity of the Sensiva formulations, are shown in Table 9.

TABLE 9

Group	<i>Staphylococcus aureus</i> CFU/patch
0.12% BZK + 0.5% PXE + 0.05% CHG + 1.0% Sensiva	30
0.12% BZK + 0.5% PXE + 0.3% PHMB + 1.0% Sensiva	20
Prevacare	1.3×10^4
Gel Base (control)	1.1×10^4
Control	1.2×10^5

EXAMPLE 8: AQUEOUS SENSIVA FORMULATION

[0040] For the experiments to be described below, the following aqueous base was used:

hydroxy methyl propyl cellulose (K100M)	0.5 percent
Kytamer PC (Chitisan)	0.15 percent
Ucare JR-400	0.1 percent
Incroquat Behenyl TMS	1.0 percent
Crodamol NM	1.6 percent
Acetulan	2.0 percent

Cremerol HMG	1.0 percent
stearyl alcohol	2.0 percent
allantoin	0.25 percent
Germall Plus	0.3 percent
dimethicone	1.0 percent (volume/volume)

and then water was added to bring to volume up to 100 percent. Various antimicrobials were added to this aqueous base, and then tested according to the protocol set forth in Example 1. The results are shown in Table 10.

TABLE 10

Group	<i>Staphylococcus aureus</i> (CFU/tube)
aqueous base (control)	5.0×10^8
0.12 % BZK	2.0×10^8
1.0 % PXE	1.0×10^8
0.5 % PXE	3.4×10^8
1.0 % Sensiva	5.0×10^8
0.05% CHG	2.5×10^8
0.3% PHMB	1.0×10^7
1% PXE + 1% Sensiva	1.0×10^8
0.05% CHG + 1% Sensiva	5.0×10^6
0.05% CHG + 1% PXE	1.0×10^8
0.12% BZK + 1% Sensiva	2.5×10^6
0.12% BZK + 1% PXE	1.2×10^7
0.12% BZK + 1% PXE + 1% Sensiva	4.0×10^4
0.12% BZK + 0.5% PXE + 0.05% CHG	2.0×10^5
0.12% BZK + 0.5% PXE + 0.05% CHG + 0.3% PHMB	2.7×10^4
0.12% BZK + 0.5% PXE + 0.05% CHG + 1% Sensiva	0
0.12% BZK + 0.5% PXE + 0.3% PHMB + 1% Sensiva	0
0.12% BZK + 0.5% PXE + 0.05% CHG + 0.3% PHMB + 1% Sensiva	0
negative control (no base/no agent)	8.0×10^8

The foregoing experiments indicate that the potentiation of the antimicrobial activity of agents by Sensiva occurs in aqueous solution, in addition to the results observed using alcoholic gels. A



combination of BZK, biguanide (CHG or PHMB), PXE and Sensiva achieved complete kill of test bacteria within 15 seconds.

EXAMPLE 9: SUSTAINED ACTIVITY OF AQUEOUS FORMULATIONS

[0041] Various combinations of antimicrobials were incorporated in an aqueous base, as set forth in Example 8, and then tested for sustained activity on leather patches using the protocol set forth in Example 7. The results, which demonstrate enhanced sustained activity in the presence of Sensiva, are shown in Table 11.

TABLE 11

Group	<i>Staphylococcus aureus</i> (CFU/patch)
0.12% BZK + 0.5% PXE + 0.05% CHG + 0.3% PHMB	2.0×10^4
0.12% BZK + 0.5% PXE + 0.05%CHG + 0.3% PHMB + 1% Sensiva	0
Aqueous Base (control)	5.0×10^5
Negative Control (no agent/no base)	5.4×10^5

EXAMPLE 10: ALCOHOL GELS CONTAINING SENSIVA AND ZINC ANTI-IRRITANTS

[0042] In individuals whose skin is sensitive to alcohol or antiseptics, the use of antimicrobial alcoholic gels can be irritating, and may cause dermatitis. It has been found that certain zinc salts, selected from the group of zinc gluconate, zinc oxide and zinc stearate, can provide an anti-irritant effect (*see* United States Patent No. 5,965,610 by Modak et al., issued October 12, 1999 and United States Patent No. 5,985,918 by Modak et al., issued November 16, 1999). In alcohol gel formulations containing Sensiva, zinc compounds were added in irritation-preventing quantities and their antimicrobial effectiveness was tested. The formulation was as follows:

zinc gluconate	2.0 percent
ethanol	63.5 percent (volume/volume)
Kytamer PC (Chitisan)	0.1 percent
Ucare JR 400 (polyquaternium 10)	0.08 percent
Germall Plus	0.3 percent
Crodamol MM	0.9 percent
Acetulan	0.5 percent
Cremerol HMG	1.0 percent
Incroquat	1.5 percent
Polawax A-31	2.0 percent
hydroxy methyl propyl cellulose (K100M)	0.4 percent
zinc stearate	3.5 percent
allantoin	0.2 percent
dimethicone	0.5 percent (volume/volume)
propylene glycol	1.5 percent (volume/volume)
glycerin	1.0 percent (volume/volume)
Sensiva	1.5 percent (volume/volume)
PXE	1.0 percent
BZK	0.12 percent
PHMB	0.3 percent

and water was added to 100% (approx. 18 ml/100 ml formulation). The resulting formulation is referred to as a "cream".

[0043] To test for rapid antimicrobial activity, 0.8 ml of the above cream formulation was mixed with 0.1 ml of 10^9 /ml CFU of test organisms and 0.1 ml bovine adult serum. After 15 seconds, this mixture was diluted 1000-fold with LTSB drug-inactivating media and 0.5 ml of the resulting solution was subcultured on a TSA plate. The resulting plates were incubated for 24 hours at 37°C and bacterial counts per tube were determined. To test for sustained antimicrobial activity, the method set forth in Example 7, using leather patches, was employed. The results of testing for rapid and sustained antimicrobial activities are shown in Table 12.

TABLE 12

Formulation	Rapid Activity (CFU/tube)	Sustained Activity (CFU/patch)
Zn Gluconate 2% + Zn Stearate 3.5% + Sensiva 1.5% + PXE 1% + BZK 0.12% + PHMB 0.3% -containing cream*	0	40
Prevacare	0	9.2×10^3
Cream Without Antimicrobials **	2.8×10^5	8.6×10^3
Control	6.5×10^8	2.3×10^5

* as comprised in the formulation set forth above in this example section.

** the formulation set forth above, omitting Sensiva, PXE, BZK and PHMB

EXAMPLE 11: ANTISEPTIC ALCOHOL GEL FORMULATION CONTAINING ZINC SALTS

[0044] The following gel formulation has only a small amount of zinc salts. It was tested for rapid antimicrobial activity against *Staphylococcus aureus*, *Pseudomonas aeruginosa*, and *Escherichia coli* using the protocol set forth in Example 10. The results, which indicate that the formulation has activity against gram positive (*Staphylococcus aureus*) as well as gram negative (*Pseudomonas aeruginosa* and *Escherichia coli*) are shown in Table 13.

ethyl alcohol	63.5 percent (volume/volume)
zinc gluconate	0.8 percent

zinc oxide	0.25 percent
hydroxy methyl propyl cellulose (K100M)	0.4 percent
Glucam P20	1.0 percent (volume/volume)
Glucam P20 distearate	1.5 percent (volume/volume)
Ucare JR400	0.15 percent
silicone (DC 1403)	1.5 percent (volume/volume)
Germall Plus	0.25 percent
PHMB	0.3 percent
PXE	1.5 percent
BZK	0.12 percent
Sensiva	1.5 percent

with water added to 100 percent (approx. 27.2 ml/100 ml).

TABLE 13

Formulation	<i>S. aureus</i> CFU/tube	<i>P. aeruginosa</i> CFU/tube	<i>E. coli</i> CFU/tube
Zn gluconate 0.8% + Zn oxide 0.2 % + PHMB 0.3% + PXE 1.5% + BZK 0.12% + Sensiva 1.5% gel*	0	1.0×10^3	0
Prevacare	0	ND	ND
Alcohol Gel Without Antimicrobials **	3.2×10^5	5.0×10^7	1.0×10^7
Control	8.0×10^8	5.0×10^8	6.5×10^8

* gel formulation set forth above in this example section.

** gel formulation set forth above, lacking PHMB, PXE, BZK and Sensiva.

EXAMPLE 12: FOAMING ANTIMICROBIAL GEL

[0045] The following alcoholic foam formulation was prepared and tested for rapid antimicrobial activity according to the method set forth in Example 10, using *Staphylococcus aureus* as the test organism. The results are shown in Table 14. If more foaming is desired, a surfactant, such as lauroyl ethylenediamine triacetic acid sodium salt (0.5-2.0%) may be added to the following formulation.

zinc gluconate	0.25 percent
zinc acetate	0.25 percent
ethanol	65.0 percent (volume/volume)
Polyquaternium 22	2.0 percent
Pluronic Gel (F-87)	0.075 percent (volume/volume)
BZK	0.12 percent
CHG	0.05%
PXE	1.0 percent
Sensiva	1.0 percent (volume/volume)

with water added to 100 percent (approx. 30.25 ml/100 ml).

TABLE 14

Formulation	<i>S. aureus</i> CFU/tube
BZK 0.12% + CHG 0.05% + PXE 1.0% + Sensiva 1.0% foam (<i>supra</i>)	0
Above Foam Without BZK, CHG, PXE or Sensiva	2.0×10^5
Control	3.9×10^8

[0046] Various publications are cited herein, the contents of which are hereby incorporated herein in their entireties by reference.